



DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration  
Rockville, MD 20857

December 19, 2005

Noam Hadas  
Scientific Laboratory Products LTD  
62 Anilevitch St.  
Tel-Aviv 67060

Dear Mr. Hadas:

Your petition requesting the Food and Drug Administration to change the classification of EEG electrodes currently classified as class II and requiring 510K approvals, was received by this office on 12/15/2005. It was assigned docket number 2005P-0213/CCP1 and it was filed on 12/15/2005. Please refer to this docket number in future correspondence on this subject with the Agency.

Please note that the acceptance of the petition for filing is a procedural matter in that it in no way reflects an agency decision on the substantive merits of the petition.

Sincerely,

Jennie C. Butler, Director  
Division of Dockets Management  
Office of Management Programs  
Office of Management

2005P-0213

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